

1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17
Sample Identification	y-Nitrocamptounecin (g)	Lipoid E80 (g)	Mannitol (g)	Trehalose (g)	Water (g)	Aq. Na Acetate* (g)	Batch size (g)	Diluent**	Dilution Factor	y-randocampioniccini Concentration, mg/g	Lipoid E80 Concentration, mg/g	Mannitol Concentration, mg/g	Trehalose Concentration, mg/g	Н	Mean Size (m)	Size: 99.9% (m)
I-A	0.252	1.051	2.74		41.5	5.0	50.6	MAN**	2.5	1.99	8.3	55.0		5.6	1.93	6.52
1-B	0.253	1.001	2.75		41.1	5.0	50.1	MAN	2.5	2.02	8.0	55.0		5.7	1.02	2.47
1-C	0.250	2.001	2.76		40.0	5.0	50.0	MAN	2.5	2.00	16.0	55.0		5.8	0.96	2.44
1-D	0.259	2.510		6.0	36.3	36.3 5.0 50.0										
1-Ea	0.250	5.000		6.0	33.8	5.0	50.1	TRE	2.5	2.00	40.0		121.3	6.0	0.07	0.22
1-Eb							-	TRE	2.5	2.00	40.0		241.3	6.0	0.07	0.20
1-Ec								TRE	2.5	2.00	40.0		361.3	6.0	0.27	2.00
1-F	1.256	25.1		15.0	71.5	12.5	125.4	TRE	5.0	2.00	40.0		240.5	5.0	1.29	2.80
1-V		16.0		9.6	46.4	8.0	80.0	TRE	5.0		4.0		239.7	4.8	0.07	0.01
,	Aq.	Na Ace	tate:									odium hydrox				
•	Dil	uent			Aqueous solution containing mannitol (MAN) or trehalose (TRE) and sodium acetate in											
					sufficient quantity to give the final concentration of sodium acetate of 2 mM and that of other											
					ingredients as shown in columns 11-14 of Table 1.											
***	Me	an Size			Volu	me we	ighted	mean p	article	diame	ter (D _{4,3})	in micromet	ers determ	ined by	a Malv	ern
						-		plus ap								
***	Siz	e:99.9%	•									this volume is apparatus.	weighted	particle	diamet	er as





St	ability of an	aqueous suspension	formulation of 9-nitrocamptothecin
	store	ed at 4°C, 25°C, and	d 40°C for up to 170 days.
Storage		veighted particle	
Temperature	diamete	r, micrometers	Appearance
and Duration	Mean	99.9 percentile	
Initial	1.29	2.80	Homogeneous yellow suspension, crystalline particles were observed in optical microscope under polarized light with a size distribution consistent with the measured size.
Stored at 4°C for 170 days	1.27	3.00	Small amounts of sediments were observed in the vial that were easily resuspendible to a
Stored at 25°C for 170 days	1.20	2.91	homogeneous yellow suspension. Crystalline particles were observed in optical microscopic
Stored at 40°C for 170 days	1.31	4.78	examination under polarized light with a size distribution consistent with the measured size. No agglomerates were found





	Initial	1.	Stress Co	ondition	•	
	Particle Size		Storage at 20°C	Storage at 40°C	4-40°C Cycling	Shaking
Test Duration	Day 0	Day18	Day18	Day18	Cycle3	Day3
Mean (volume weighted)	0.20 µm	0.19 μm	0.18 µm	0.17 μm	0.19 µm	0.20 µm
99.9 Percentile	0.34 μm	0.34 μm	0.31 μm	0.31 μm	0.33 µm	0.33 μm

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FIGURE 4

Protocol Design For First Melanoma Xenograft Study

		TreatmentRegimen 1	Regimen1		
dnoio	=	Agent	mg/kg	Route	Schedule
-	10	NoTreatment	n/a	n/a	
2	01	IDD-P(1:3dilution)	n/a	.≥	5/2/5
3	0_	IDD-D(nodilution)	n/a	.≥	5/2/5
4	10	D5Wwith3%DMA	n/a	od	Day1,4,8,11
5	01	CAMPTOSAR	100	.₽	QWKx3
9	10	HYCAMTIN	10	ġ.	Q4Dx4
7	10	DTIC	150	di	QDx5
∞	10	9NC-IDD-P	3	vi	5/2/5
6	10	9NC-IDD-P	1.5	.≥	5/2/5
10	10	9NC-IDD-D	2	Ņ	5/2/5
=	10	9NC-IDD-D	1	.≥	5/2/5
12	10	9NC-D5W-3%DMA	4	od	Day1,4,8,11
13	10	9NC-D5W-3%DMA	2	bo	Day1,4,8,11



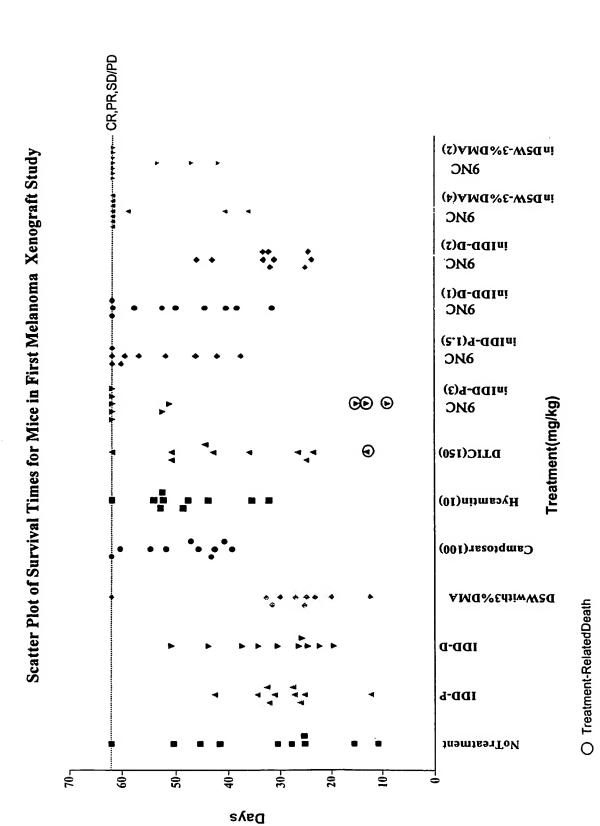


Treatment Response Summary For First Melanoma Xenograft Study

		TreatmentRegimen	Regime	g		MDSto2.0g	Max.%BW	#De	#Death 1	d C	40,7	Car Car
Group	=	Agent	mg/kg	Route	Schedule	±SEM(n)	Loss;Day	TR	NTR	#CK	#1.IK	#3D/I'U
-	10	NoTreatment	n/a	n/a		30.3 ± 4.4 (9)	*	0	0	0	0	1
2	10	IDD-P(1:3dilution)	n/a	iv	5/2/5	29.2 ± 2.5 (10)		0	0	0	0	0
3	10	IDD-D(nodilution)	n/a	iv	5/2/5	31.6 ± 3.2 (10)	-0.4%;Day27	0	0	0	0	0
4	10	D5Wwith3%DMA	n/a	bo	Day 1, 4, 8, 11	26.0 ± 2.3 (8)		0	_		0	0
5	10	CAMPTOSAR	100	di	QWKx3	47.3 ± 2.3 (9)		0	0	_	0	0
9	10	HYCAMTIN	10	ţ	Q4Dx4	46.7 ± 2.6 (9)		0	0	-	0	0
7	10	DTIC	150	qi	, QDx5	37.6 ± 4.0 (8)	-5.2%;Day5	-	0	_	0	0
8	01	9NC-IDD-P	3	iv	5/2/5	52.0 ± 0.6 (2)	-13.1%;Day13		0	_	3	
6	10	9NC-IDD-P	1.5	Ņ	5/2/5	50.7 ± 3.4 (7)	-2.2%;Day5	0	0	2	0	
01	01	9NC-IDD-D	2	Ņ	5/2/5	47.2 ± 3.6 (8)	•	0	0	_	_	0
11	10	9NC-IDD-D	-	iv	5/2/5	32.6 ± 2.3 (10)	***	0	0	0	0	0
12	01	9NC-D5W-3%DMA	4	od	Day 1, 4, 8, 11	45.3 ± 7.0 (3)	•	0	0	4	0	3
13	10	9NC-D5W-3%DMA	2	od	Day1,4,8,11	47.6 ± 3.4 (3)	1	0	0	4	~	2

 ${}^{a}\#\mathrm{Dcath}.TR(TreatmentRelated); NTR(Non-TreatmentRelated)$

FIGURE 6



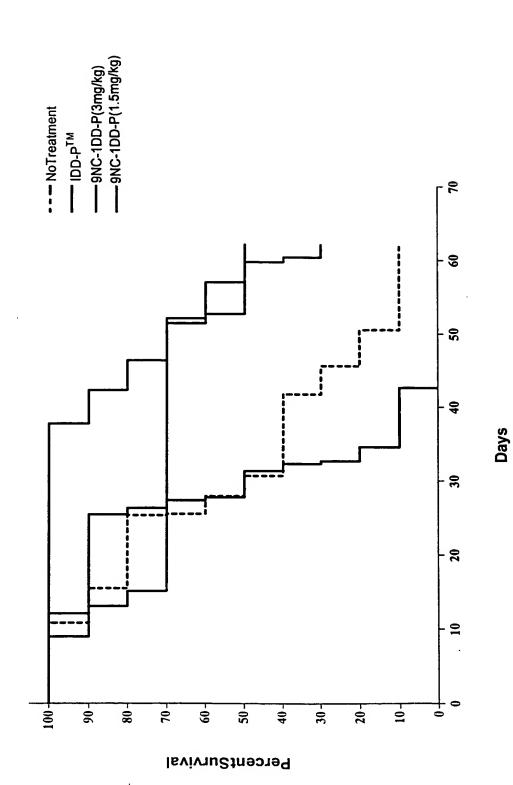


Figure 7



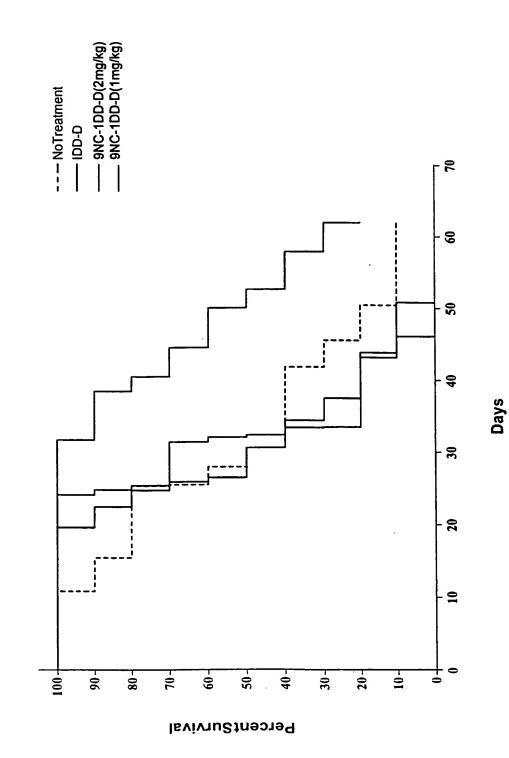
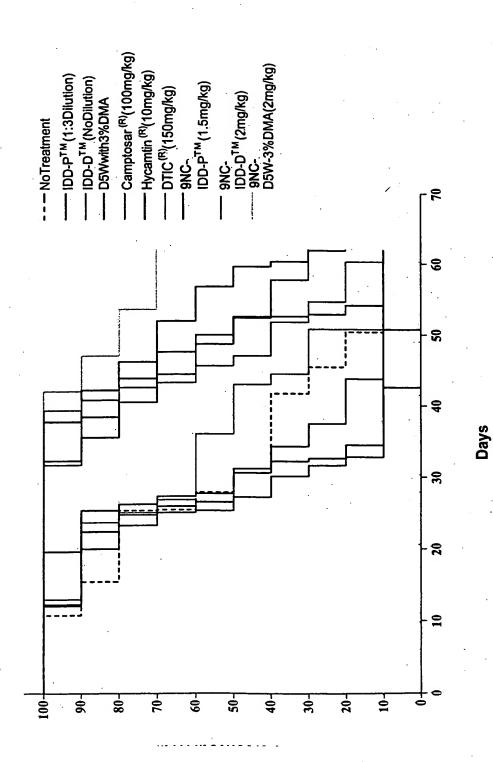
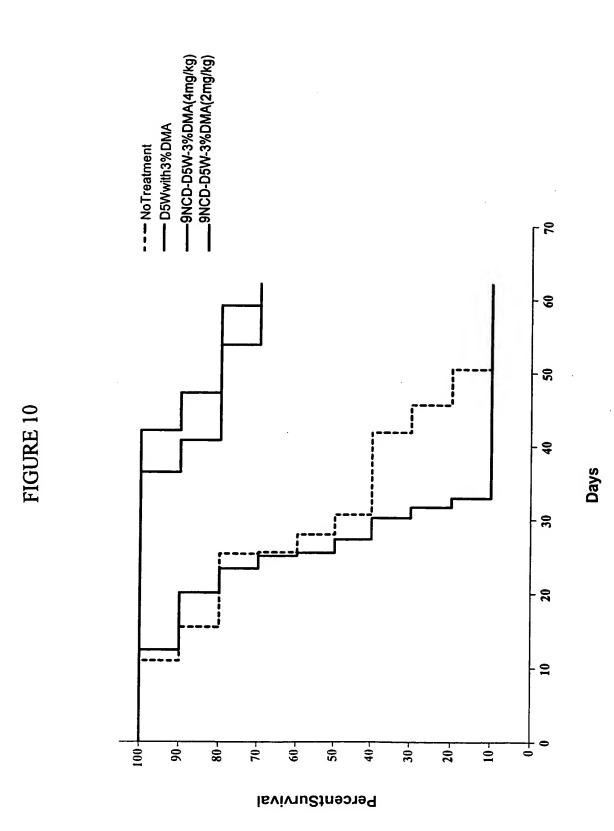


Figure 8













5/2/5 + 5 daily dose, 2 days rest, 5 daily Qwkx3 one dose per day at four day intervals QDx5 one dose per day for 5 days

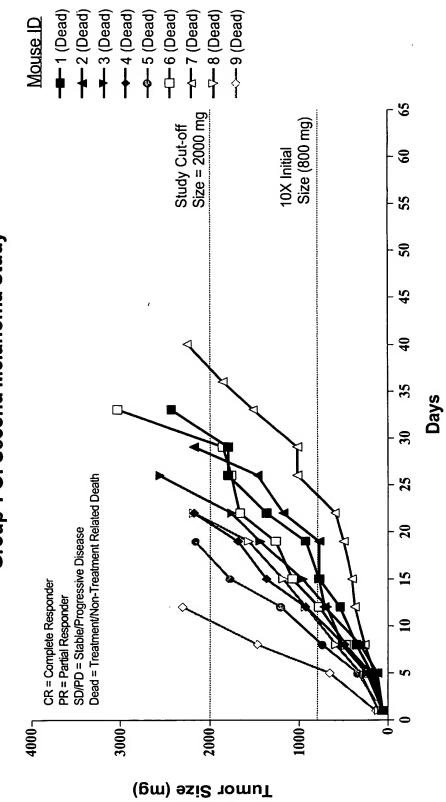
01/01	\$000.0>>>	1.5	9:45	9NC 2 mg/kg oral
01/01	\$000.0>>>	8.£	1.42	9NC 4 mg/kg oral
				l mg/kg iv
01/01	11.0	p .1	26.1	
				vi gy∕kg iv
01/01	\$000.0>>>	1.5	6.14	9AC in IDD-D
				vi ga/kg iv
01/01	\$000.0>>>	Τ.ε	2.24	9AC in IDD-P
				3 mg/kg iv
<i>L</i> /01	\$000.0>>>	1.5	2.95	9NC-in IDD-P
				(cxQQ) qi
6/0 I	90.0	9.₽	5.15	DTIC 150 mg/kg
				(4xQ4Q) qi
01/01	2000.0	2.9	6.65	Hycamtin 10 mgkg
				ip (Qwkx3)
01/01	<u></u> ₽ 000.0	2.5	7.65	Camptosar 100mg/kg
				(\$/7/\$)
6/01	21.0	0.1	15.2	AMG driw Wed
				(5/2/5) vi
01/01	84.0	4.2	18.0	IDD-D vehicle
			_	(2/2/5) vi
10/10	25.0	9.£	£.7.I	IDD-P vehicle
01/01		0.5	20.5	No treatment
day 62				
ot gaivivme				
reaching 2 g or	treatment			
n\n lsitinl	on sv q	+_ SEM_	Mean (days)	Treatment

EICHE 11



FIGURE 12A

Group 1 Of Second Melanoma Study





Study Cut-off Size = 2000 mg **Group 2 of Second Melanoma Study FIGURE 12 B** SD/PD = Stable/Progressive Disease

4000-

3000-

→ 3 (SD/PD) → 4 (SD/PD) → 5

9 -

62

9

55

20

45

30

25

20

15

10

Days

10X Initial Size (800 mg)

1000 -

2000-

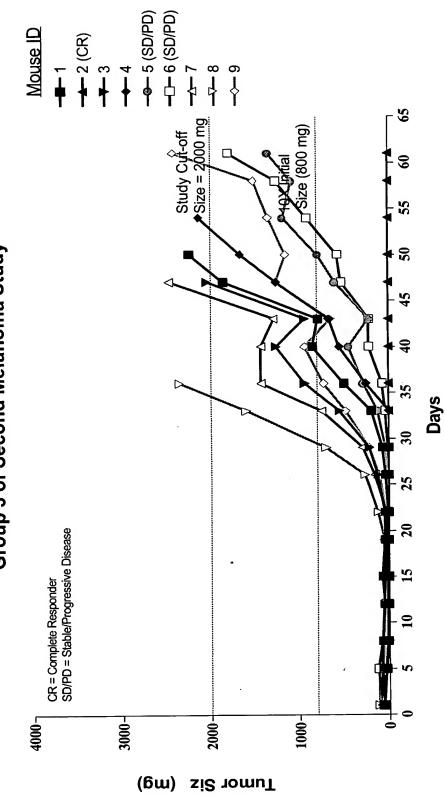
Tumor Size (mg)

Mouse ID



FIGURE 12C

Group 3 of Second Melanoma Study





→ 3 (CR) → 4 (SD/PD) → 5 (CR) → 1 ← 6 (SD/PD) → 7 (CR) **-** 1 (SD/PD) **-** 2 (SD/PD) —7—8 (SD/PD) Mouse ID 65 Study Cut-off Size = 2000 mg Size (800 mg) Group 5 of Second Melanoma Study Days CR = Complete Responder SD/PD = Stable/Progressive Disease 4000 € 1000 -3000-2000 Tumor Size (mg)



FIGURE 12D



FIGURE 13

MX-1 Human Breast Cancer Xenograft Study

Treatment	Schedule	Mean Days to 10 X	S.E.M	P vs. no treatment	# of mice at start/ # mice reaching 10x
No treatment		17.4	2.23		10/10
IDD-P vehicle i.v.	5/2/5	16.5	1.1	n.s	10/10
9NC in IDDP 2.5 mg/kg i.v.	5/2/5	53.0	0.0	<<0.05	10/10
9NC in IDDP 1.75 mg/kg i.v.	5/2/5	53.0	0.0	<<0.05	10/10
9NC in IDDP 1.25 mg/kg i.v.	5/2/5	47.5	2.1	<<0.05	10/10.
Camptosar 100 mg/kg i.p	QWK x 3	53.0	0.0	<<0.05	10/10
Hycamtin 10 mg/kg i.p.	Q4D x 4	53.0	0.0	<<0.05	10/10



FIGURE 14

Pan 1- Human Pancreatic Cancer Xenograft Study

Treatment	Schedule	Mean Days to 10 X	S.E.M	P vs. no treatment	# of mice at start/ # mice reaching 10x
No treatment		19.5	1.6		10/10
IDD-P vehicle i.v.	5/2/5	20.6	1.3	n.s	9/9
9NC in IDDP 2.5 mg/kg i.v.	5/2/5	34.3	2.0	<<0.05	10/10
9NC in IDDP 1.75 mg/kg i.v.	5/2/5	25.7	1.3	<0.01	10/10
9NC in IDDP 1.25 mg/kg i.v.	5/2/5	24.6	1.0	=.01	10/10.
Camptosar 100 mg/kg i.p	QWK x 3	30.5	3.9	<0.05	10/10
Hycamtin 10 mg/kg i.p.	Q4D x 4	30.6	1.5	<<0.05	10/10



FIGURE 15

HT-29 Human Colon Cancer Xenograft Study

Treatment	Schedule	Mean Days to 10 X	S.E.M	P vs. no treatment	# of mice at start/ # mice reaching 10x
No treatment		26.9	2.0		8/8
IDD-P vehicle i.v.	5/2/5	29.4	1.6	n.s	8/8
9NC in IDDP 2.5 mg/kg i.v.	5/2/5	34.0	1.8	<0.05	8/8
9NC in IDDP 1.75 mg/kg i.v.	5/2/5	34.5	2.0	<0.05	9/9
9NC in IDDP 1.25 mg/kg i.v.	5/2/5	38.1	3.6	<0.05	9/9.
Camptosar 100 mg/kg i.p	QWK x 3	35.7	2.2	<0.01	9/9
Hycamtin 10 mg/kg i.p.	Q4D x 4	34.4	1.5	<0.01	9/9



FIGURE 16
SKMES Human Lung Cancer Xenograft Study

Treatment	Schedule	Mean Days to 10 X	S.E.M	P vs. no treatment	# of mice at start/ # mice reaching 10x
No treatment		11.7	0.8		10/10
IDD-P vehicle i.v.	5/2/5	14.6	1.0	0.03	10/10
9NC in IDDP 2.5 mg/kg i.v.	5/2/5	27.3	1.6	<<0.05	10/10
9NC in IDDP 1.75 mg/kg i.v.	5/2/5	29.4	2.2	<<0.05	10/10
9NC in IDDP 1.25 mg/kg i.v.	5/2/5	35.2	5.7	<0.05	10/10.
Camptosar 100 mg/kg i.p	QWK x 3	35.2	4.4	<<0.05	10/10
Hycamtin 10 mg/kg i.p.	Q4D x 4	33.6	3.6	<<0.05	10/10